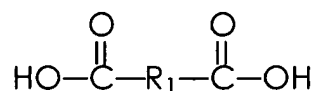


1. A process for preparing a tetra-amide which comprises carrying out a condensation reaction between a diacid, a monoacid, and a diisocyanate, thereby forming a tetra-amide.

2. A process according to claim 1 wherein the diacid is of the formula



wherein R₁ is an alkylene group, an arylene group, an arylalkylene group, or an alkylarylene group.

3. A process according to claim 2 wherein R₁ is an unsubstituted alkylene group, an unsubstituted arylene group, an unsubstituted arylalkylene group, or an unsubstituted alkylarylene group.

4. A process according to claim 2 wherein R₁ is a substituted alkylene group, a substituted arylene group, a substituted arylalkylene group, or a substituted alkylarylene group.

5. A process according to claim 2 wherein R₁ is an alkylene group having hetero atoms therein, an arylene group having hetero atoms therein, an arylalkylene group having hetero atoms therein, or an alkylarylene group having hetero atoms therein, provided that no hetero atoms are directly bonded to either of the carboxylic acid groups.

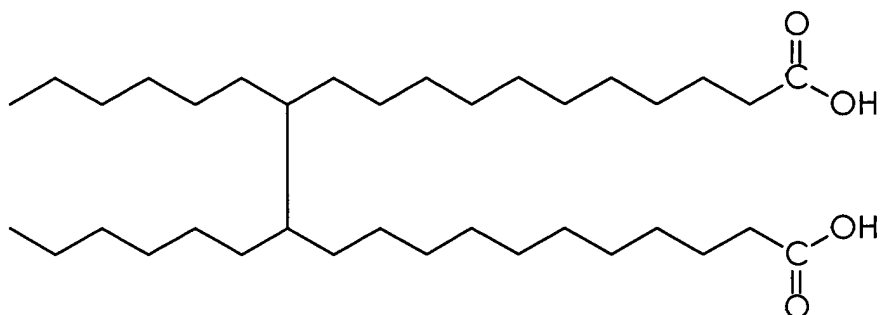
6. A process according to claim 2 wherein R_1 is an alkylene group having no hetero atoms therein, an arylene group having no hetero atoms therein, an arylalkylene group having no hetero atoms therein, or an alkylarylene group having no hetero atoms therein.

7. A process according to claim 2 wherein R_1 is a branched alkylene group having at least about 34 carbon atoms.

8. A process according to claim 2 wherein R_1 is a branched alkylene group having about 34 carbon atoms and which may include unsaturations and cyclic groups.

9. A process according to claim 2 wherein R_1 is a branched alkylene group of the formula $C_{34}H_{62+n}$ wherein n is an integer of 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10.

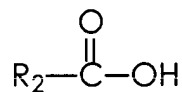
10. A process according to claim 1 wherein the diacid is of the formula



11. A process according to claim 1 wherein the diacid is malonic acid, methyl malonic acid, ethyl malonic acid, butyl malonic acid, dimethyl malonic acid, diethyl malonic acid, succinic acid, methyl succinic acid, dimethyl succinic acid, 2-ethyl-2-methyl succinic acid, 2,3-dimethyl succinic acid, glutaric acid, 2-methyl glutaric acid, 3-methyl glutaric acid, 2,2-dimethyl glutaric acid, 3,3-dimethyl glutaric acid, adipic acid, 3-methyl adipic acid, 3-tert-butyl adipic acid, pimelic acid, suberic acid, azelaic acid, sebacic acid, 1,11-undecanedicarboxylic acid, undecanedioic acid, 1,10-decanedicarboxylic acid, 1,12-dodecanedicarboxylic acid, hexadecanedioic acid, docosanedioic acid, tetracosanedioic acid, itaconic acid, maleic acid, fumaric acid, citraconic acid, mesaconic acid, glutaconic acid, β -hydromuconic acid, traumatic acid, muconic acid, aconitic acid, chlorosuccinic acid, bromosuccinic acid, 2,3-dibromosuccinic acid, tetrafluorosuccinic acid, hexafluoroglutaric acid, perfluoroadipic acid, perfluorosuberic acid, 3-chlorododecanedioic acid, dibromomaleic acid, diglycolic acid, 3,6-dioxaoctanedioic acid, thiodiglycolic acid, 3,3'-thiodipropionic acid, 1,3-acetonedicarboxylic acid, 3-oxoadipic acid, 4-ketopimelic acid, 5-oxoazelaic acid, chelidonic acid, 1,2-cyclopentanedicarboxylic acid, 3,3-tetramethyleneglutaric acid, camphoric acid, cyclohexylsuccinic acid, 1,1-cyclohexanediacetic acid, 1,2-cyclohexanedicarboxylic acid, 1,3-cyclohexanedicarboxylic acid, 1,4-cyclohexanedicarboxylic acid, 1,3-adamantanedicarboxylic acid, 1,3-adamantanediacetic acid, 5-norbornene-2,3-dicarboxylic acid, 1,4,5,6,7,7-hexachloro-5-norbornene-2,3-dicarboxylic acid, phenylsuccinic acid, 3-phenylglutaric acid, 1,2-phenylenediacetic acid, 1,2-phenylenedioxydiacetic acid, homophthalic acid, 1,3-phenylenediacetic acid, 4-

carboxyphenoxyacetic acid, 1,4-phenylenediacetic acid, 1,4-phenylenedipropionic acid, 2-carboxycinnamic acid, 1,4-phenylenediacrylic acid, 2-carboxybenzenepropanoic acid, 4,4'-(hexafluoroisopropylidene)bis(benzoic acid), 4,4'-oxybis(benzoic acid), phthalic acid, isophthalic acid, terephthalic acid, 3-fluorophthalic acid, 2-methoxyisophthalic acid, 3-nitrophthalic acid, 4-methylphthalic acid, 2-bromoterephthalic acid, 4-bromoisophthalic acid, 4-nitrophthalic acid, nitroterephthalic acid, 5-tert-butylisophthalic acid, 5-octadecyloxyisophthalic acid, 5-nitroisophthalic acid, 4,5-dichlorophthalic acid, tetrafluoroterephthalic acid, tetrafluoroisophthalic acid, tetrafluorophthalic acid, diphenic acid, 4,4'-biphenyldicarboxylic acid, 4-[4-(2-carboxybenzoyl)phenyl]butyric acid, 1,4-naphthalenedicarboxylic acid, 2,3-naphthalenedicarboxylic acid, 2,6-naphthalenedicarboxylic acid, 2,7-di-tert-butyl-9,9-dimethyl-4,5-xanthenedicarboxylic acid, phenylmalonic acid, benzylmalonic acid, or mixtures thereof.

12. A process according to claim 1 wherein the monoacid is of the formula



wherein R₂ is an alkyl group, an aryl group, an arylalkyl group, or an alkylaryl group.

13. A process according to claim 12 wherein R₂ is an unsubstituted alkyl group, an unsubstituted aryl group, an unsubstituted arylalkyl group, or an unsubstituted alkylaryl group.

14. A process according to claim 12 wherein R₂ is a substituted alkyl group, a substituted aryl group, a substituted arylalkyl group, or a substituted alkylaryl group.

15. A process according to claim 12 wherein R₂ is an alkyl group having hetero atoms therein, an aryl group having hetero atoms therein, an arylalkyl group having hetero atoms therein, or an alkylaryl group having hetero atoms therein, provided that no hetero atoms are directly bonded to the carboxylic acid group.

16. A process according to claim 2 wherein R₁ is an alkylene group having no hetero atoms therein, an arylene group having no hetero atoms therein, an arylalkylene group having no hetero atoms therein, or an alkylarylene group having no hetero atoms therein.

17. A process according to claim 1 wherein the monoacid is acetic acid, propionic acid, butyric acid, valeric acid, hexanoic acid, heptanoic acid, octanoic acid, nonanoic acid, decanoic acid, undecanoic acid, lauric acid, tridecanoic acid, myristic acid, pentadecanoic acid, palmitic acid, heptadecanoic acid, stearic acid, nonadecanoic acid, eicosanoic acid, heneicosanoic acid, docosanoic acid, tricosanoic acid, tetracosanoic acid, hexacosanoic acid, heptacosanoic acid, octacosanoic acid, triacontanoic acid, isobutyric acid, 2-ethylbutyric acid, trimethylacetic acid, 2-methylbutyric acid, isovaleric acid, 2,2-dimethylbutyric acid, tert-butylacetic acid, 2-methylvaleric acid, 2-propylpentanoic acid, 3-methylvaleric acid, 4-methylvaleric acid, 2-methylhexanoic acid, 2-ethylhexanoic acid, acrylic acid, methacrylic acid, crotonic acid, vinylacetic acid, tiglic acid, 3,3-dimethylacrylic acid, 2-pentenoic acid, 4-pentenoic acid, 2-methyl-2-pentenoic acid, 2,2-dimethyl-4-pentenoic acid, 2-hexenoic acid, 3-hexenoic acid, 2-ethyl-2-hexenoic acid, 6-heptenoic acid, 2-octenoic acid, citronellic acid, undecylenic acid, myristoleic acid, palmitoleic acid, oleic acid, elaidic acid, 11-eicosenoic acid, erucic acid, nervonic acid, chloroacetic acid, bromoacetic acid, iodoacetic acid, difluoroacetic acid, dichloroacetic acid, dibromoacetic acid, trifluoroacetic acid, chlorodifluoroacetic acid, trichloroacetic acid, tribromoacetic acid, 2-chloropropionic acid, 3-chloropropionic acid, 2-bromopropionic acid, 3-bromopropionic acid, 2-iodopropionic acid, 3-iodopropionic acid, 2,2-dichloropropionic acid, 2,3-dibromopropionic acid, pentafluoropropionic acid, 2-bromo-2-methylpropionic acid, 3-bromo-2-(bromomethyl)-propionic acid, 3-chloropivalic acid, 3,3-dichloropivalic acid, 4-chlorobutyric acid, 2-bromobutyric acid, 4-bromobutyric acid, heptafluorobutyric acid, 2-

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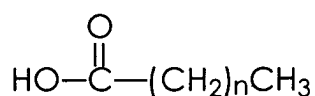
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acid, 2-methoxy-4-(methylthio)benzoic acid, 5-methyl-2-nitrobenzoic acid, 4-methyl-3-nitrobenzoic acid, 3-methyl-4-nitrobenzoic acid, 2-nitro- α,α,α -trifluoro-p-toluic acid, 2-fluoro-5-nitrobenzoic acid, 4-chloro-2-nitrobenzoic acid, 2-chloro-4-nitrobenzoic acid, 4-fluoro-3-nitrobenzoic acid, 4-chloro-3-nitrobenzoic acid, 5-chloro-2-nitrobenzoic acid, 2-chloro-5-nitrobenzoic acid, 2-bromo-5-nitrobenzoic acid, 4-(bromomethyl)-3-nitrobenzoic acid, 2-methoxy-4-nitrobenzoic acid, 4-methoxy-3-nitrobenzoic acid, 3-methoxy-4-nitrobenzoic acid, 5-methoxy-2-nitrobenzoic acid, 2,4-dinitrobenzoic acid, 3,5-dimethylbenzoic acid, 3,5-di-tert-butylbenzoic acid, 3,5-difluorobenzoic acid, 3,5-bis(trifluoromethyl)benzoic acid, 3,5-dichlorobenzoic acid, 3,5-dibromobenzoic acid, 3-bromo-5-iodobenzoic acid, 3,5-dimethoxybenzoic acid, 3,5-dinitrobenzoic acid, 2,3,4-trifluorobenzoic acid, 2,3,6-trifluorobenzoic acid, 2,4,6-trimethylbenzoic acid, 2,4,6-trifluorobenzoic acid, 3,4,5-trifluorobenzoic acid, 2,4,6-trichlorobenzoic acid, 2,3,5-trichlorobenzoic acid, 2,3,5-triiodobenzoic acid, 2-bromo-4,5-dimethoxybenzoic acid, 3,4,5-trimethoxybenzoic acid, 3,4,5-triethoxybenzoic acid, 4,5-dimethoxy-2-nitrobenzoic acid, 3,5-dinitro-o-toluic acid, 3,5-dinitro-p-toluic acid, 2-chloro-3,5-dinitrobenzoic acid, 4-chloro-3,5-dinitrobenzoic acid, 2,5-dichloro-3-nitrobenzoic acid, 2,6-dichloro-3-nitrobenzoic acid, 2,3,4-trimethoxybenzoic acid, 2,4,5-trifluorobenzoic acid, 2-chloro-4,5-difluorobenzoic acid, 2,4-dichloro-5-fluorobenzoic acid, 2,4,5-trimethoxybenzoic acid, 2,3,4,5-tetrafluorobenzoic acid, 2,3,5,6-tetrafluorobenzoic acid, 2,4-dichloro-3,5-dinitrobenzoic acid, 2,3,5,6-tetrafluoro-p-toluic acid, 4-bromo-2,3,5,6-tetrafluorobenzoic acid, pentafluorobenzoic acid, 2-biphenylcarboxylic acid, 4'-(trifluoromethyl)-2-biphenylcarboxylic acid, 4-biphenylcarboxylic acid, 4'-ethyl-4-biphenylcarboxylic acid, 4'-octyloxy-

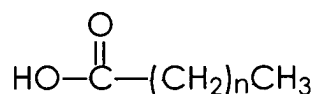
4-biphenylcarboxylic acid, α -phenyl-o-toluic acid, 2-bibenzylcarboxylic acid, 2,3,4,5,6-pentafluorophenoxyacetic acid, 2-phenoxybenzoic acid, 3-phenoxybenzoic acid, 2-benzoylbenzoic acid, 3-benzoylbenzoic acid, 4-benzoylbenzoic acid, 2-(4-fluorobenzoyl)benzoic acid, 2-(4-chlorobenzoyl)benzoic acid, 2-(4-chloro-3-nitrobenzoyl)benzoic acid, 1-naphthoic acid, 2-naphthoic acid, 4-fluoro-1-naphthoic acid, 2-ethoxy-1-naphthoic acid, 1,8-naphthalaldehydic acid, 2-biphenylenecarboxylic acid, γ -oxo-5-acenaphthenebutyric acid, 9-fluorenicarboxylic acid, 1-fluorenicarboxylic acid, 4-fluorenicarboxylic acid, 9-fluorenone-1-carboxylic acid, 9-fluorenone-2-carboxylic acid, 9-fluorenone-4-carboxylic acid, 7-nitro-4-fluorenicarboxylic acid, chromone-2-carboxylic acid, 9-anthracenecarboxylic acid, anthraquinone-2-carboxylic acid, xanthene-9-carboxylic acid, 1-pyrenecarboxylic acid, or mixtures thereof.

18. A process according to claim 1 wherein the monoacid is of the formula



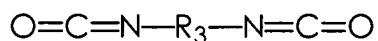
wherein n has an average value of about 36.

19. A process according to claim 1 wherein the monoacid is of the formula



wherein n has an average value of about 46.

20. A process according to claim 1 wherein the diisocyanate is of the formula



wherein R_3 is an alkylene group, an arylene group, an arylalkylene group, or an alkylarylene group.

21. A process according to claim 20 wherein R_3 is an unsubstituted alkylene group, an unsubstituted arylene group, an unsubstituted arylalkylene group, or an unsubstituted alkylarylene group.

22. A process according to claim 20 wherein R_3 is a substituted alkylene group, a substituted arylene group, a substituted arylalkylene group, or a substituted alkylarylene group.

23. A process according to claim 20 wherein R_3 is an alkylene group having hetero atoms therein, an arylene group having hetero atoms therein, an arylalkylene group having hetero atoms therein, or an alkylarylene group having hetero atoms therein, provided that no hetero atoms are directly bonded to either of the isocyanate groups.

24. A process according to claim 20 wherein R_3 is an alkylene group having no hetero atoms therein, an arylene group having no hetero atoms therein, an arylalkylene group having no hetero atoms therein, or an alkylarylene group having no hetero atoms therein.

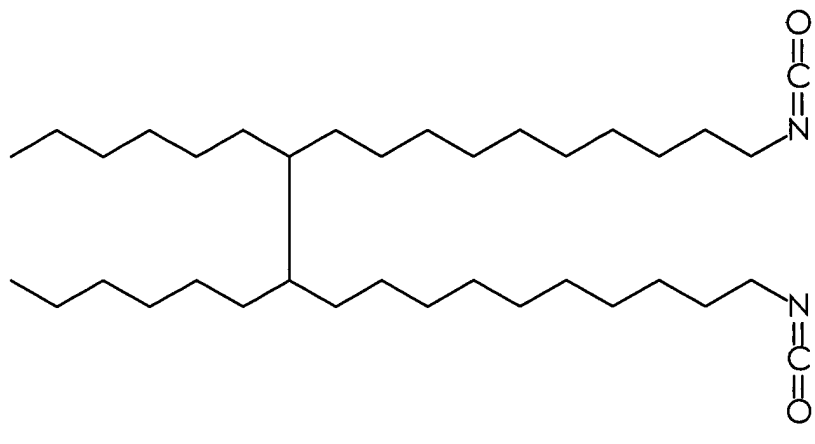
25. A process according to claim 20 wherein R_3 is a branched alkylene group having about 34 carbon atoms.

26. A process according to claim 20 wherein R_3 is an alkylene group of the formula $C_{34}H_{62+n}$ wherein n is an integer of 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10.

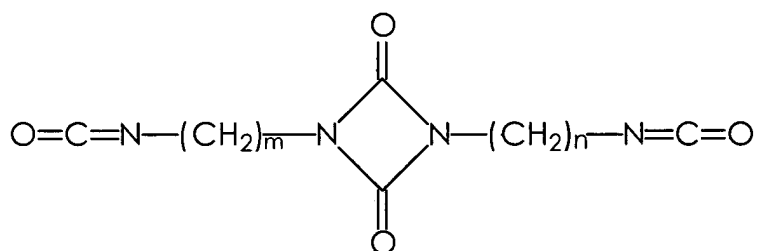
27. A process according to claim 1 wherein the diisocyanate is 1,4-diisocyanatobutane, 1,6-diisocyanatohexane, 1,8-diisocyanatooctane, 1,12-diisocyanatododecane, trimethylhexamethylene diisocyanate, 1,5-diisocyanato-2-methylpentane, cyclohexylene diisocyanate, bis(isocyanatomethane) cyclohexane, 4,4'-methylenebis(cyclohexyl isocyanate), isophorone diisocyanate, phenylene diisocyanate, bis(isocyanatomethyl)benzene, bis(1-isocyanato-1-methylethyl)benzene, toluene diisocyanate, diphenylmethane-4,4'-diisocyanate, hydrogenated diphenylmethane-4,4'-diisocyanate, tetramethylxylene diisocyanate, naphthylene-1,5-diisocyanate, 3,3'-dimethoxy-4,4'-biphenyldiisocyanate, 3,3'-dimethyl-4,4'-bimethyl-4,4'-biphenyldiisocyanate, 4,4'-biphenyldiisocyanate, tetramethylene xylene diisocyanate, 4,4'-methylenebis(2,6-diethylphenyl isocyanate), 1-chloromethyl-2,4-diisocyanatobenzene, 4,4'-oxybis(phenyl isocyanate), or mixtures thereof.

28. A process according to claim 1 wherein the diisocyanate is isophorone diisocyanate or 1,6-diisocyanatohexane.

29. A process according to claim 1 wherein the diisocyanate is of the formula



30. A process according to claim 1 wherein the diisocyanate is of the formula

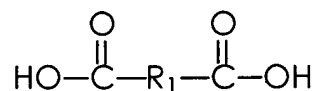


wherein m and n are each, independently of the others, integers representing the number of repeat -CH₂- units.

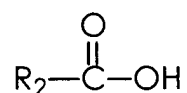
31. A process according to claim 30 wherein m and n are each 6.

32. A process according to claim 1 wherein the tetra-
amide has a total number of carbon atoms of at least about 50.

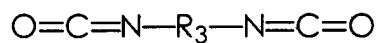
33. A process according to claim 1 wherein the diacid is of the formula



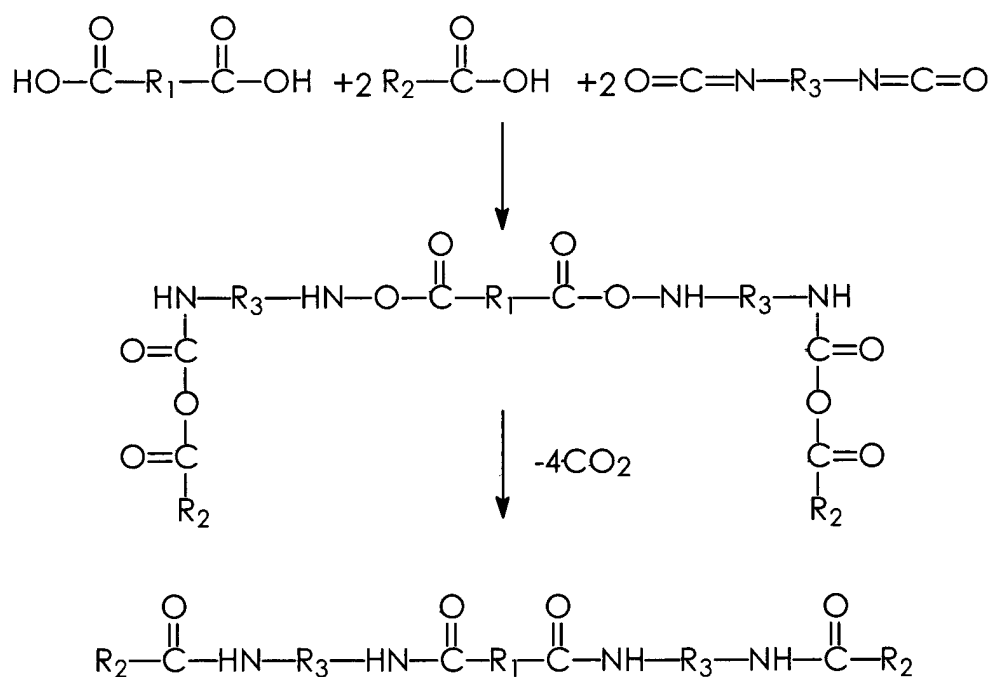
wherein R_1 is an alkylene group, an arylene group, an arylalkylene group, or an alkylarylene group, the monoacid is of the formula



wherein R_2 is an alkyl group, an aryl group, an arylalkyl group, or an alkylaryl group, the diisocyanate is of the formula



wherein R_3 is an alkylene group, an arylene group, an arylalkylene group, or an alkylarylene group, and the reaction proceeds as follows:



34. A process according to claim 1 wherein the diacid and the monoacid are first admixed prior to addition of the diisocyanate reactant.

35. A process according to claim 1 wherein the diacid and the monoacid are first admixed and heated prior to addition of the diisocyanate reactant.

36. A process according to claim 1 wherein the diacid, the monoacid, and a polyethylene wax are first admixed and heated prior to addition of the diisocyanate reactant.

37. A process according to claim 1 wherein the reaction takes place in an inert atmosphere.

38. A process according to claim 1 wherein the reaction mixture is heated to a temperature of at least about 60°C.

39. A process according to claim 1 wherein the reaction mixture is heated to a temperature of at least about 80°C.

40. A process according to claim 1 wherein the reaction mixture is heated to a temperature of at least about 120°C.

41. A process according to claim 1 wherein the reaction mixture is heated to a temperature of no more than about 400°C.

42. A process according to claim 1 wherein the reaction mixture is heated to a temperature of no more than about 300°C.

43. A process according to claim 1 wherein the reaction mixture is heated to a temperature of no more than about 200°C.

44. A process according to claim 1 wherein the reaction is carried out for a period of at least about 10 minutes.

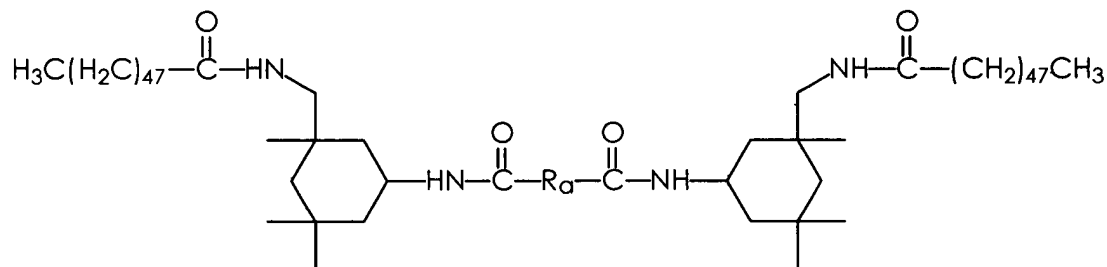
45. A process according to claim 1 wherein the reaction is carried out for a period of at least about 30 minutes.

46. A process according to claim 1 wherein the reaction is carried out for a period of at least about 60 minutes.

47. A process according to claim 1 wherein the monoacid, the diacid, and the diisocyanate are present in relative amounts such that the ratio of total number of COOH groups to total number of NCO groups is at least about 1:1.

48. A process according to claim 1 wherein a catalyst is added to the reaction mixture.

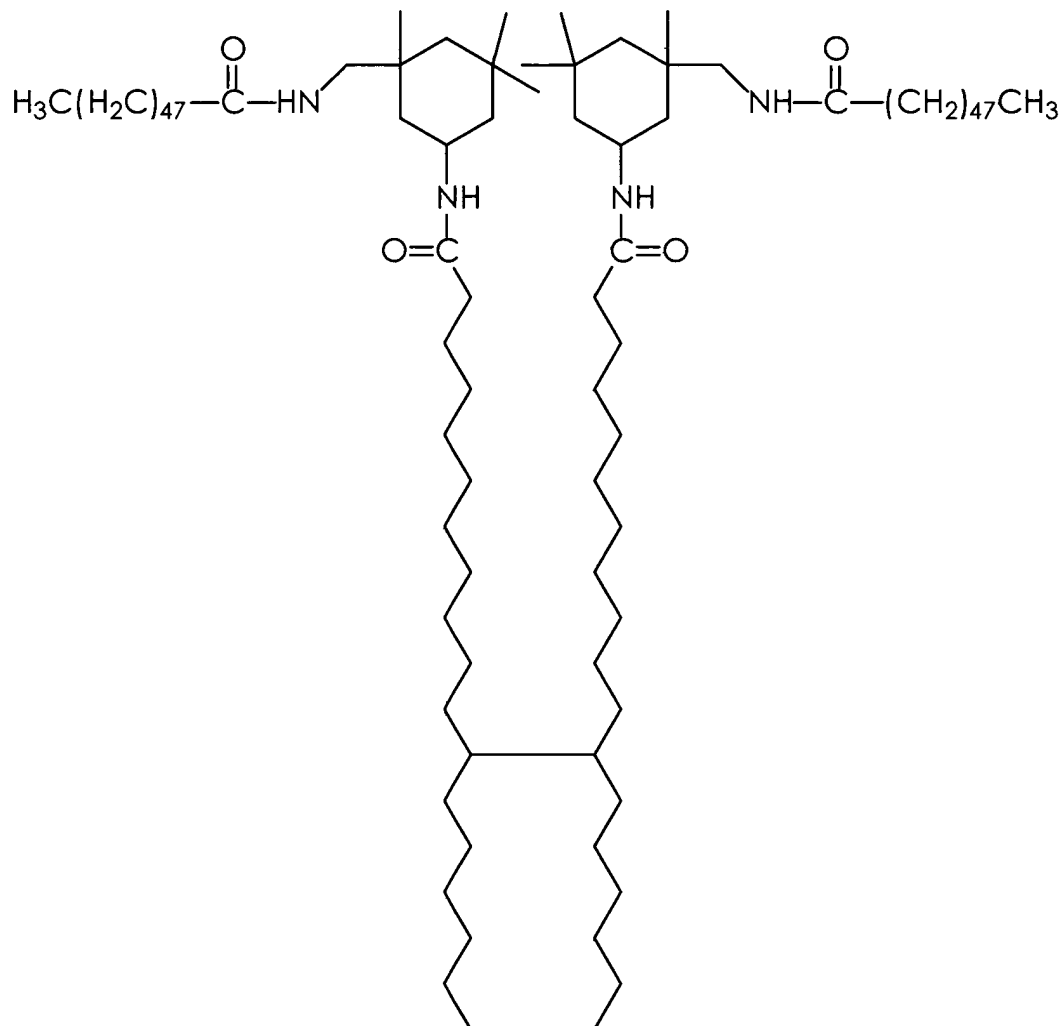
49. A process according to claim 1 wherein the tetra-amide product is of the formula



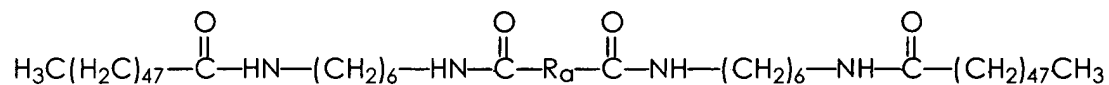
wherein R_a is a branched alkylene group having about 34 carbon atoms and which may include unsaturations and cyclic groups.

50. A process according to claim 49 wherein R_a is a group of the formula $C_{34}H_{62+n}$ wherein n is an integer of 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10.

51. A process according to claim 1 wherein the tetra-
amide product is of the formula



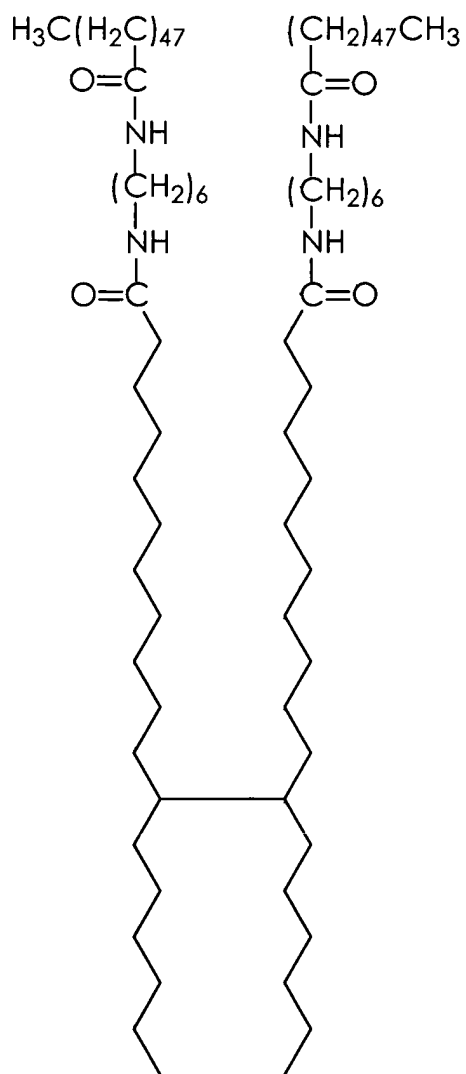
52. A process according to claim 1 wherein the tetra-
amide product is of the formula



wherein R_a is a branched alkylene group having about 34 carbon atoms and which may include unsaturations and cyclic groups.

53. A process according to claim 52 wherein R_a is a group of the formula $C_{34}H_{62+n}$ wherein n is an integer of 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, or 10.

54. A process according to claim 1 wherein the tetra-amide product is of the formula



55. A process for preparing a phase change ink composition which comprises (a) carrying out a condensation reaction between a diacid, a monoacid, and a diisocyanate, thereby forming a tetra-amide; and (b) admixing the tetra-amide thus formed with a colorant, thereby forming a phase change ink.

56. A process according to claim 55 wherein the tetra-amide and the colorant are further admixed with a polyethylene wax.

57. A process according to claim 55 wherein the tetra-amide and the colorant are further admixed with a monoamide.

58. A process according to claim 55 wherein the tetra-amide and the colorant are further admixed with a urethane resin.

59. A process according to claim 55 wherein the tetra-amide is present in the ink in an amount of at least about 10 percent by weight of the ink and in an amount of no more than about 32 percent by weight of the ink, and wherein the tetra-amide and the colorant are further admixed with (i) a polyethylene wax, present in the ink in an amount of at least about 25 percent by weight of the ink and in an amount of no more than about 60 percent by weight of the ink; (ii) a stearyl stearamide wax, present in the ink in an amount of at least about 8 percent by weight of the ink and in an amount of no more than about 32 percent by weight of the ink; (iii) a urethane resin derived from the reaction of two equivalents of hydroabietyl alcohol and one equivalent of isophorone diisocyanate, present in the ink in an amount of at least about 6 percent by weight of the ink and in an amount of no more than about 16 percent by weight of the ink; (iv) a urethane resin that is the adduct of three equivalents of stearyl isocyanate and a glycerol-based alcohol, present in the ink in an amount of at least about 2 percent by weight of the ink and in an amount of no more than about 13 percent by weight of the ink; and (v) an antioxidant, present in the ink in an amount of at least about 0.01 percent by weight of the ink and in an amount of no more than about 1 percent by weight of the ink.

60. A process according to claim 55 wherein the tetra-amide is present in the ink in an amount of at least about 13 percent by weight of the ink and in an amount of no more than about 27 percent by weight of the ink, and wherein the tetra-amide and the colorant are further admixed with (i) a polyethylene wax, present in the ink in an amount of at least about 30 percent by weight of the ink and in an amount of no more than about 53 percent by weight of the ink; (ii) a stearyl stearamide wax, present in the ink in an amount of at least about 10 percent by weight of the ink and in an amount of no more than about 28 percent by weight of the ink; (iii) a urethane resin derived from the reaction of two equivalents of hydroabietyl alcohol and one equivalent of isophorone diisocyanate, present in the ink in an amount of at least about 8 percent by weight of the ink and in an amount of no more than about 14 percent by weight of the ink; (iv) a urethane resin that is the adduct of three equivalents of stearyl isocyanate and a glycerol-based alcohol, present in the ink in an amount of at least about 3 percent by weight of the ink and in an amount of no more than about 10 percent by weight of the ink; and (v) an antioxidant, present in the ink in an amount of at least about 0.05 percent by weight of the ink and in an amount of no more than about 0.5 percent by weight of the ink.

61. A process according to claim 55 wherein the tetra-amide is present in the ink in an amount of at least about 16 percent by weight of the ink and in an amount of no more than about 22 percent by weight of the ink, and wherein the tetra-amide and the colorant are further admixed with (i) a polyethylene wax, present in the ink in an amount of at least about 37 percent by weight of the ink and in an amount of no more than about 48 percent by weight of the ink; (ii) a stearyl stearamide wax, present in the ink in an amount of at least about 12 percent by weight of the ink and in an amount of no more than about 25 percent by weight of the ink; (iii) a urethane resin derived from the reaction of two equivalents of hydroabietyl alcohol and one equivalent of isophorone diisocyanate, present in the ink in an amount of at least about 10 percent by weight of the ink and in an amount of no more than about 12 percent by weight of the ink; (iv) a urethane resin that is the adduct of three equivalents of stearyl isocyanate and a glycerol-based alcohol, present in the ink in an amount of at least about 4.5 percent by weight of the ink and in an amount of no more than about 7.5 percent by weight of the ink; and (v) an antioxidant, present in the ink in an amount of at least about 0.1 percent by weight of the ink and in an amount of no more than about 0.3 percent by weight of the ink.

62. A process according to claim 55 wherein the colorant is present in the ink in an amount of at least about 0.1 percent by weight of the ink.

63. A process according to claim 55 wherein the colorant is present in the ink in an amount of no more than about 10 percent by weight of the ink.

64. A process according to claim 55 wherein the ink has a melting point of no lower than about 50°C.

65. A process according to claim 55 wherein the ink has a melting point of no lower than about 70°C.

66. A process according to claim 55 wherein the ink has a melting point of no lower than about 80°C.

67. A process according to claim 55 wherein the ink has a melting point of no higher than about 160°C.

68. A process according to claim 55 wherein the ink has a melting point of no higher than about 140°C.

69. A process according to claim 55 wherein the ink has a melting point of no higher than about 100°C.

70. A process according to claim 55 wherein the ink has a viscosity at jetting temperatures of no more than about 30 centipoise.

71. A process according to claim 55 wherein the ink has a viscosity at jetting temperatures of no more than about 20 centipoise.

72. A process according to claim 55 wherein the ink has a viscosity at jetting temperatures of no more than about 15 centipoise.

73. A process according to claim 55 wherein the ink has a viscosity at jetting temperatures of no less than about 2 centipoise.

74. A process according to claim 55 wherein the ink has a viscosity at jetting temperatures of no less than about 5 centipoise.

75. A process according to claim 55 wherein the ink has a viscosity at jetting temperatures of no less than about 7 centipoise.